

VA Natalizumab (Tysabri®) Clinical Monitoring Program
Initial Registry Information

Date of Evaluation: ____/____/____

VAMC Healthcare Provider: _____

VAMC Provider Phone #: _____ Email: _____

Name of VA Facility: _____

VAMC Location (City): _____ State: _____ Facility/Station #: _____

☐ Check here if transferring natalizumab (Tysabri®) treatment from another VA.

Name of Patient (first, last name): _____

Date of Birth: ____/____/____

Patient's Four Digit VA Code: ____

Complete at the initial visit prior to starting natalizumab (Tysabri®)

1. Sex:

☐ Male
 ☐ Female
2. Race:

☐ American Indian or Alaskan Native
☐ Asian
☐ Black or African American

☐ Native Hawaiian or Pacific Islander
☐ White
☐ Other: _____
3. Ethnicity:

☐ Hispanic or Latino
 ☐ Non-Hispanic or Latino
4. Year of onset of initial MS symptoms: _____
5. Number of relapses over the past year (prior to starting natalizumab(Tysabri®)): _____
6. Indicate the MS disease modifying therapies used in the past:
 - a. Interferon-beta:
 - i. Avonex®: total months on therapy: _____
 - ii. Betaseron®: total months on therapy: _____
 - iii. Rebif®: total months on therapy: _____
 - b. Glatiramer acetate: Copaxone®: total months on therapy: _____
 - c. Mitoxantrone: Novantrone®: total months on therapy: _____
 - d. Chemotherapy/Other: _____ (name medication) total months on therapy: _____
7. Indication(s) for Natalizumab (Tysabri®), check all that apply:

☐ Side effects from interferon-beta
☐ Side effects from glatiramer acetate
☐ Other: _____

☐ Inadequate response despite interferon-beta therapy
☐ Inadequate response despite glatiramer acetate therapy
8. MS Disease Subtype:

☐ Relapsing-remitting
 ☐ Secondary-progressive with relapses
 ☐ Progressive-relapsing
9. MS Disability at time of evaluation:
 - a. Expanded Disability Status Scale (Kurtzke J, et al *Neurology* 1983;13:1444) *check box*:

<input type="checkbox"/> 0	<input type="checkbox"/> 2.0	<input type="checkbox"/> 3.0	<input type="checkbox"/> 4.0	<input type="checkbox"/> 5.0	<input type="checkbox"/> 6.0	<input type="checkbox"/> 7.0	<input type="checkbox"/> 8.0	<input type="checkbox"/> 9.0
<input type="checkbox"/> 1.0	<input type="checkbox"/> 2.5	<input type="checkbox"/> 3.5	<input type="checkbox"/> 4.5	<input type="checkbox"/> 5.5	<input type="checkbox"/> 6.5	<input type="checkbox"/> 7.5	<input type="checkbox"/> 8.5	<input type="checkbox"/> 9.5
<input type="checkbox"/> 1.5								
 - or**
 - b. Provider Determined Disease Steps (Hohol M, et al *Neurology* 1995;45:251) *check box*:

<input type="checkbox"/> 0-Normal	<input type="checkbox"/> 4-Late Cane
<input type="checkbox"/> 1-Mild Disability	<input type="checkbox"/> 5-Bilateral Support
<input type="checkbox"/> 2-Moderate Disability	<input type="checkbox"/> 6-Wheelchair
<input type="checkbox"/> 3-Early Cane	
10. Pretreatment Brain MRI by CMSC Protocol (www.va.gov/ms) completed: ☐ Date: ________ (mo/yr)

Please FAX this form to: Alicia Sloan, MPH, LICSW, Natalizumab Clinical Monitoring Program Coordinator

MS Center of Excellence-West, FAX: **206-277-4827**, VOICE: 206-277-3593

Questions? Email Alicia.Sloan@va.gov

Visit the MSCoE website at www.va.gov/ms Note: This is not a research protocol.